525 Rec'd PCT/PTO 24 OCT 2000

FORM PTO-1390 (REV 12-29-99) | U S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE | ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371

Mo-5946/WW-5504

U S APPLICATION NO (If known, see 37 CFR 1.5)

To Be Assigned 09/673944

INTERNATIONAL APPLICATION NO.	INTERNATIONAL FILING DATE	PRIORITY DATE CLAIMED
PCT/EP99/02553	April 16, 1999	April 28, 1998

TITLE OF INVENTION Method for Producing Transparent, Colored Cellulose Sleeves

APPLICANT(S) FOR DO/EO/US BLUMENBERG, Klaus-Dieter and NEUSCHULZ, Willi

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. [X] This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. [ ] This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. [X] This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. [X] A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
5. [X] A copy of the International Application as filed (35 U.S.C. 371(c)(2))
   a. [X] is transmitted herewith (required only if not transmitted by the International Bureau).
   b. [ ] has been transmitted by the International Bureau.
   c. [ ] is not required, as the application was filed in the United States Receiving Office (RO/US).
6. [X] A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. [ ] Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
   a. [ ] are transmitted herewith (required only if not transmitted by the International Bureau).
   b. [ ] have been transmitted by the International Bureau.
   c. [ ] have not been made; however, the time limit for making such amendments has NOT expired.
   d. [ ] have not been made and will not be made.
8. [ ] A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. [X] An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. [ ] A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern document(s) or information included:

11. [ ] An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. [X] An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. [X] A FIRST preliminary amendment.
    [ ] A SECOND or SUBSEQUENT preliminary amendment.
14. [ ] A substitute specification.
15. [ ] A change of power of attorney and/or address letter.
16. [X] Other items or information:

Abstract
Form PTO 1449 w/references

528 Rec'd PCT/PTO 24 OCT 2000

U.S. APPLICATION NO (If known, see 37 CFR 1.5)	INTERNATIONAL APPLICATION NO	ATTORNEY'S DOCKET NUMBER
To Be Assigned 09/673744	PCT/EP99/02553	Mo-5946/WW-5504

17. [X] The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

	CALCULATIONS PTO USE ONLY
Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO **$970.00**	
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO **$840.00**	
International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO **$690.00**	
International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) **$670.00**	
International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4) **$96.00**	
ENTER APPROPRIATE BASIC FEE AMOUNT =	$ 860.00
Surcharge of **$130.00** for furnishing the oath or declaration later than [ ] 20 [ ] 30 months from the earliest claimed priority date (37 CFR 1.492(e)).	$ 0.00

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	
Total claims	9 - 20 =	0	X **$18.00**	$ 0.00
Independent claims	1 - 3 =	0	X **$78.00**	$ 0.00
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ **$260.00**	$ 0.00
TOTAL OF ABOVE CALCULATIONS =				$ 860.00
Reduction of 1/2 for filing by small entity, if applicable. A Small Entity Statement must also by filed (Note 37 CFR 1.9, 1.27, 1.28).				$ 0.00
SUBTOTAL =				$ 860.00
Processing fee of **$130.00** for furnishing the English translation later than [ ]20 [ ] 30 months from the earliest claimed priority date (37 CFR 1.492(f)). +				$ 0.00
TOTAL NATIONAL FEE =				$ 860.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). **$40.00** per property +				$ 40.00
TOTAL FEES ENCLOSED =				$ 900.00
			Amount to be refunded:	$
			charged:	$

a. [ ] A check in the amount of $_____ to cover the above fees is enclosed.

b. [X] Please charge my Deposit Account No. 13-3848 in the amount of $ 900.00 to cover the above fees. A duplicate copy of this sheet is enclosed.

c. [X] The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 13-3848. A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO

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00157
PATENT TRADEMARK OFFICE

SIGNATURE

Joseph C. Gil
NAME

26,602
REGISTRATION NUMBER

09/673944
528 Rec'd PCT/PTO 24 OCT 2000

PATENT APPLICATION
Mo-5946
WW-5504

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICATION OF	)
	) PCT/EP99/02553
KLAUS-DIETER BLUMENBERG ET AL	)
	)
SERIAL NUMBER: TO BE ASSIGNED	)
	)
FILED: HEREWITH	)
	)
TITLE: METHOD FOR PRODUCING TRANSPARENT, COLORED CELLULOSE SLEEVES	)

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Prior to the examination of the subject patent application, kindly amend the enclosed, English language translation thereof as follows:

IN THE SPECIFICATION:

Please delete the title of the translated document as it appears in page 1 thereof and insert therefor a revised title reading: --Method for Producing Transparent, Colored Cellulose Sleeves--;

And in page 15, a page containing an abstract, delete the indicated title and insert therefor: --Method for Producing Transparent, Colored Cellulose Sleeves--.

"Express Mail" mailing label number EF080092405US
Date of Deposit October 24, 2000

I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above and is addressed to the Assistant Commissioner of Patents and Trademarks, Washington, D.C. 20231

Donna J. Vearch
(Name of person mailing paper or fee)

Signature of person mailing paper or fee)

IN THE CLAIMS:

In Claim 3, line 1, please delete "claim 1 or 2," and insert --Claim 1--, therefor.

In Claim 4, line 1, please delete "claims 1 to 3," and insert --Claim 1--, therefor.

In Claim 5, line 1, please delete "claims 1 to 4," and insert --Claim 1--, therefor.

In Claim 6, line 1, please delete "claims 1 to 5," and insert --Claim 1--, therefor.

In Claim 7, line 3, please delete "one of claims 1 to 6," and insert --Claim 1--, therefor.

Cancel Claim 9.

Add Claim 10 that reads:

-- 10. A method of using the tubular wrapping of Claim 7 comprising preparing a synthetic casing for a member selected from the group consisting of raw sausage, sausage for boiling and sausage for cooking.--

REMARKS

The amendment seeks to render the translated application in better conformance with U.S. practice. A page containing the amended Abstract of the Disclosure is enclosed.

An early examination on the merits is respectfully solicited.

Respectfully submitted,

KLAUS-DIETER BLUMENBERG
WILLI NEUSCHULZ

By _____
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METHOD FOR PRODUCING TRANSPARENT, COLORED CELLULOSE SLEEVES

ABSTRACT OF THE DISCLOSURE

Process for producing dyes, tubular food wrappings from non-woven fabric coated with regenerated cellulose, characterized in that an alkaline dye liquor containing at least one dye which has been previously converted into an alkali-soluble form by chemical reduction and which can be converted into its insoluble form by oxidation is admixed to the viscose solution used for the production of the layer of regenerated cellulose, a tubular non-woven fabric is coated with the mixture of viscose solution and dye liquor, the viscose is coagulated and regenerated to form cellulose hydrate gel and the dye distributed in the viscose is reconverted into its insoluble form by oxidation; tubular food wrappings produced by this process and their use as synthetic casings for sausages.

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Date of Deposit October 24, 2000
I hereby certify that this paper or fee is being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 on the date indicated above and is addressed to the Assistant Commissioner of Patents and Trademarks, Washington, D.C. 20231
Donna J. Veatch
(Name of person mailing paper or fee)
(Signature of person mailing paper or fee)

09/673944
Received PCT/PTO 24 OCT 2000

Process for producing transparent dyed cellulose wrappings

This invention relates to a process for producing tubular dyed wrappings for food, in particular skins made of cellulose fibre, having high transparency and evenness of the dye, and also to the tubular food wrappings produced according to the invention and the use thereof.

Tubular food wrappings are used on a large scale in the production of a multitude of meat products. These food wrappings are generally thin-walled tubes of various diameters and in many cases are produced in known manner from regenerated cellulose. Cellulose wrappings can also be produced in the form of skins of cellulose fibre containing embedded fibrous material.

The appearance of sausages produced using skins of cellulose fibre is an important factor in appealing to the consumer, and a multitude of products are cured in order to obtain a characteristic brown coloration. It is also usual to dye skins of cellulose fibre, for which purpose coloured pigments and dyes are used. Particularly in the case of raw sausages in which the sausage meat has a coarse-grained structure, such as, for example, salami, it is desirable that the lumps of meat and fat should show clearly through the sausage casing in order to achieve an appearance which will promote sale. In the conventional pigment dyeing of skins made of cellulose fibre, the transparency of the sausage casings is greatly reduced owing to the diffuse refraction of light on the relatively large particles of dye.

The dyeing of cellulose films and in particular cellulose fibres with so-called vat dyes is known. The vat dyes are compounds having an indigoid or anthraquinonoid structure and are insoluble in water. They require a special dyeing process, the essential features of which have long been known. In this process, with the use of reducing agents, such as sodium hydrogen sulfite, sodium dithionite, sodium hydroxymethanesulfinate or sodium borohydride, the vat dye is first of all converted into its completely alkali-soluble leuco form. The cellulose material to be dyed is

dipped into or passed through this dye liquor, which is referred to as vat. The high affinity for cellulose of the leuco dyes brings about a high rate of dyeing at the surface of the cellulose material, which can lead to unevennesses in the dye in cases where mixtures of dyes in which the leuco dyes have differing affinities for cellulose are used. There is then a diffusion of the leuco dyes into the interior of the cellulose material, the diffusion being accelerated by elevation of the temperature. After the material has been rinsed, the oxidation is carried out and the original water-insoluble dye, which adheres well to the cellulose material, is again formed from the leuco form. Examples of suitable oxidising agents are atmospheric oxygen, hydrogen peroxide, sodium perborate and potassium dichromate.

Thus US Patent No. 3,149,905 claims the dyeing and printing of cellulose-containing textile materials with vat dyes. The vat dyes used according to that patent contain sulfonamide groups. The dyeing is carried out in conventional dipping baths.

The dyeing of cellulose materials from regenerated cellulose, cellulose esters and cellulose ethers is claimed in US Patent 2,043,069. In this process, the vat dye in its undissolved form is mixed with the cellulose starting compound and the coagulation and regeneration are carried out. The regenerated cellulose material is subsequently passed through the reducing and oxidising baths in order to dissolve and fix the vat dyes.

A very similar process is claimed in Deutsche Offenlegungsschrift 2,262,611 for the production of cellulose films. Here the vat dye in its undissolved form is mixed with the viscose, the regeneration is carried out and the vat dye is subsequently dissolved and fixed in reducing and oxidising baths.

Comparable dyeing processes using vat dyes have not hitherto been applied, however, to the group of products comprising skins made of cellulose fibre.

A process for dyeing a skin of cellulose fibre in the gel state in dye liquors is claimed in US Patent 3,383,443. However, the dyes used there are not vat dyes but naphthol dyes, which are fixed with stabilised diazo compounds.

Finally, US Patent 3,293,340 describes a dyeing process which uses vat dyes for the production of wrappings from transparent dyed, regenerated cellulose. In this process, the still undissolved vat dye in the form of a pigment paste is mixed with the viscose and from this mixture is produced a tube of regenerated cellulose, which is regenerated in acid coagulating baths in conventional manner and subsequently washed. The use of an additional desulfurisation bath prior to or after the reduction step is recommended. The tube is then passed through an alkaline reducing bath, the vat dye being converted into the alkali-soluble leuco form. The reducing bath contains in addition 100 to 150 g/l sodium chloride in order to suppress the migration of the soluble leuco form out of the tube. The reconversion of the leuco form into the insoluble form is then effected by oxidation with atmospheric oxygen in a special duct, in which water is sprayed onto the tube in order to facilitate oxidation by removing the reducing agent. The tube is subsequently washed again and preferably passed through another acid bath having a concentration of 1.5 to 7.5 g sulfuric acid per litre, in order to neutralise the rest of the sodium hydroxide and thereby to enable the tube to be washed more efficiently in the subsequent repeated washing step. The conventional treatment with softeners in a glycerol bath then takes place, and finally the drying step.

The process described in US Patent 3,293,340, owing to the use of the additional processing steps involving the reducing bath, the oxidising chamber, the intermediate washing, the subsequent acid treatment and the repeated washing, is capital-intensive as compared with the conventional processes for producing fibre-reinforced cellulose wrappings. Furthermore, because of the necessity of controlling the concentration in the reducing bath and in the acid bath, it is very expensive and in addition may altogether be seen as contributing to the waste-water loading. Where a reducing bath is used, the undesirable oxidation of the reducing agent by

atmospheric oxygen cannot be avoided, owing to the large surface area in contact with the air. Considerably more reducing agent is therefore consumed than is necessary for the conversion of the dyes into the leuco form, in consequence of which the waste water is additionally polluted. In particular, the high concentration of sodium chloride in the reducing bath claimed according to the invention in the patent under discussion leads to additional waste-water loading, which is to be regarded as highly problematic in view of the environmental protection generally practised nowadays.

In the above process, moreover, the process of diffusion of the reducing agent into the regenerated tube has a decisive influence on the evenness and transparency of the dye obtained. This gives rise to difficulties in the application of the process, as skins of cellulose fibre are conventionally produced using non-woven fabric of varying thickness and varying quantities of applied viscose, depending upon the caliber. Apart from the thickness of the regenerated tube, factors which influence the diffusion rate are the temperature and concentration of the reducing bath. These influences can barely be controlled, especially when variously dyed tubes of different caliber are passed simultaneously through a reducing bath.

Accordingly, the object of the present invention was to provide a process for producing dyed, tubular food wrappings made of non-woven fabric coated with regenerated cellulose, in particular skins of cellulose fibre, having high transparency and evenness of the dye, which can be carried out at low industrial expense. In particular, the quantities of the chemicals used are to be as small as possible, so as to minimise environmental pollution.

This object is achieved by the process described in claim 1. Claims 2 to 6 give preferred embodiments of the process. The object is also achieved by the product according to claim 7, by its development according to claim 8 and by its use according to claim 9.

Surprisingly, it has been found that dyes, in particular vat dyes, which are convertible into alkali-soluble form, after chemical reduction can, in their alkali-soluble leuco form, be homogeneously mixed with viscose and with this mixture dyed tubular food wrappings made of non-woven fibre coated with regenerated cellulose, in particular skins of cellulose fibre, having high transparency and evenness of dye can be easily produced by admixing to the viscose solution used for the production of the layer of regenerated cellulose an alkaline dye liquor containing at least one dye which has been previously converted into an alkali-soluble form by chemical reduction and which can be converted into its insoluble form by oxidation; by coating a tubular non-woven fabric with the mixture of viscose solution and dye liquor; by coagulating the viscose and regenerating it to form cellulose hydrate gel and by reconverting the dye distributed in the viscose into its insoluble form by oxidation.

The dyes preferably used for the process according to the invention are dyes of the class of substances comprising the anthraquinone derivatives, in particular derivatives of anthrimidecarbazole, acylaminoanthraquinone, acridone, benzanthrone, violanthrone, isoviolanthrone, indanthrene, and derivatives of more highly condensed aromatic ring systems, preferably pyrenequinone, anthanthrone, flavanthrone, pyranthrone, perylenetetracarboxylic acid, naphthalenetetracarboxylic acid as well as indigo derivatives and thioindigo derivatives.

The dyed, tubular food wrapping produced by the process according to the invention is particularly suitable for use as synthetic casings for sausages and can in addition be provided with a barrier layer on the outer and inner surface. Food wrappings produced by the process according to the invention are used particularly preferably as synthetic casings for raw sausage, sausages for boiling or sausages for cooking.

The process according to the invention for producing a transparent, dyed fibre-reinforced cellulose wrapping preferably proceeds as follows:-

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Commercially available vat dyes in the form of coloured pastes or powders are chemically reduced in separate tanks and dissolved in sodium hydroxide solution prior to being mixed with viscose. For example, the vat dye or mixtures of different vat dyes in the form of coloured paste or powder are placed in a tank and water, or preferably an aqueous solution of one or more cellulose ethers, in particular carboxymethyl cellulose or methylene cellulose, is added thereto. Preferably types of these cellulose ethers which are of low viscosity are used, the viscosity of their 2 wt.% solutions being preferably 30 to 300 mPa·s. A freshly prepared aqueous solution of sodium dithionite or sodium sulfide and sodium hydroxide is added thereto. It is also possible to place the vat dyes in a tank and then to add the other substances simultaneously, dissolved in water. The components are mixed with a stirrer, the rate of rotation being so adjusted that no air bubbles are introduced into the dye liquor. In the course of a residence time of preferably 12 to 16 hours at room temperature, the vat dyes are completely reduced to the leuco form and are dissolved in the lye. Afterwards, the dye liquor is stirred briefly and introduced into the receiving tanks of conventional units for mixing viscose and dye. If a toning down of the dye is desired, alkali- and acid-resistant coloured pigments in a quantity of 3 to 12 wt.%, preferably 4 to 7 wt.%, based on the total quantity of dye can also be used in this process, in addition to mixtures of vat dyes. Here the addition of cellulose ethers, besides bringing about a diminished rate of uptake of atmospheric oxygen as a result of the increase in viscosity, also effects the dispersion of the insoluble coloured pigments in the dye liquor. For the chemical reduction of the vat dyes, sodium dithionite or sodium sulfide is used in a quantity of 10 to 90 wt.%, preferably 20 to 80 wt.%, based on the pure vat dyes in the dye liquor.

Depending on the solubility of the leuco forms of the vat dyes and on the required intensity of the dye, the concentration of the leuco form in the dye liquor is 1.25 to 4.4 wt.%. The concentration of the sodium dithionite in the dye liquor is preferably 0.7 to 2.0 wt.% and that of the sodium hydroxide is 0.3 to 0.9 wt.%.

The process according to the invention henceforth makes possible the precise adjustment of the quantities of chemicals to the respective vat dyes and hence the prevention of their irreversible overreduction. In particular the introduction of large excess quantities of reducing agents, which are necessary where reducing baths are used owing to oxidation by atmospheric oxygen, is omitted. The coagulation of the viscose in the course of being mixed with the dye liquor is prevented by the low concentrations of salt in the dye liquor. Agglomeration of dye in the dyed sausage casing is consequently avoided.

If relatively prolonged storage of the finally prepared dye liquor is required, the containers should be filled with it as fully as possible and closed with an air-tight seal, in order to prevent oxidation by atmospheric oxygen. The units for mixing viscose and dye are conventional devices for the continuous mixing of the dye liquor and the viscose. A fixed proportion of viscose to metered dye liquor is established by means of adjustable delivery pumps. The addition of the dye liquor to the viscose is 2 to 26 litres, preferably 3 to 9 litres, per 100 kg viscose. The receiving tanks of the units for mixing viscose and dye can be closed with an air-tight seal and the gas space above the dye liquor can be flushed with nitrogen in order to prevent the oxidation of the dye liquor by atmospheric oxygen during longer production runs. The dyed viscose is passed via piping or tubing to the viscosing nozzles conventionally used. Here the web of non-woven fabric shaped in the form of a tube, which is adjusted depending on the tube diameter to be produced, is preimpregnated and coated on one or both sides with the dyed viscose. The tube of non-woven fabric coated with dyed viscose is passed through a conventional, acid- and salt-containing system of coagulating baths, during which the tube in various places is fully immersed in the bath and also passes through vertical air gaps. The viscose is converted into regenerated cellulose by the sulfuric acid contained in the baths and at the same time the leuco form of the indanthrene dyes contained therein is reconverted by oxidation with atmospheric oxygen into the original, insoluble dye, which adheres firmly to the cellulose material. Afterwards the tube of regenerated cellulose is washed in conventional manner, passed through a softening bath and

dried with supporting air. Where the wrapping is used in the production of sausages for cooking and sausages for boiling, a barrier layer against atmospheric oxygen and water vapour may optionally be applied to the outer or inner side by known methods.

In this process claimed according to the invention, commercially available vat dyes which are convertible into an alkali-soluble form by chemical reduction can be used such as, for example, the violanthrone derivative indanthrene brilliant green (C.I. Part 1: vat green 1, C.I. Part 2: 59825), the naphthalenetetracarboxylic acid derivative indanthrene brilliant orange GR (C.I. Part 1: vat orange 7, C.I. Part 2: 71105), the anthanthrone derivative indanthrene brilliant orange RF (C.I. Part 1: vat orange 3, C.I. Part 2: 59300), the thioindigo derivative indanthrene brilliant pink R (C.I. Part 1: vat red 1, C.I. Part 2: 73360), the violanthrone derivative indanthrene blue (C.I. Part 1: vat blue 20, C.I. Part 2: 59800), the benzanthrone derivative indanthrene grey (C.I. Part 1: vat black 8, C.I. Part 2: 71000), indanthrene scarlet GG (C.I. Part 1; vat red 14, C.I. Part 2:71110), the naphthalenetetracarboxylic acid derivative indanthrene Bordeaux RR (C.I. Part 1: vat red 15, C.I. Part 2: 71100), the acylaminoanthraquinone derivative indanthrene yellow F3GC (C.I. Part 1: vat yellow 33, C.I. Part 2: 65430), the benzopyrenequinone indanthrene yellow GOK (C.I. Part 1: vat yellow 4, C.I. Part 2: 59100) and perylenetetracarboxylic acid derivatives, such as paliogenmarron (C.I. Part 1: pigment red 179, C.I. Part 2: 71130). C.I. in each case denotes the generic name and/or constitution number according to the Colour Index issued by the Society of Dyers and Colourists.

Coloured pastes or powders of these pigments are marketed, for example, under the names Suprafix pastes or Colloisol.

The invention is illustrated in more detail by the following Examples.

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Examples

Example 1

In a sealable container of 30 litres in volume, 30.04 kg of a dye liquor was prepared as follows:-

6 kg	commercially available dye paste containing approximately 22% of the dye Red 179, C.I. 71130 (6300 Vocofil Rot, Ari Chemie) and
210 g	pigment black 7, C.I. 77266, pigment preparation with the name of Novofil Schwarz BB03, pigment content approximately 35% (water-insoluble pigment)

were weighed out. A freshly prepared solution of

230 g	carboxymethyl cellulose (CRT 30 GA, Wolff Walsrode)
500 g	sodium hydroxide solution (50 wt%)
600 g	sodium dithionite

in 22.5 kg of water was added thereto.

The components were gently stirred with a paddle mixer, the rate of rotation being so adjusted that no air bubbles were introduced into the dye liquor. After a residence time of 12 hours in the sealed container, this dye liquor was mixed continuously in constant ratio with viscose, in a ratio of 3.3 l dye liquor to 100 kg viscose. A web of non-woven fabric having a width of cut corresponding to a nominal caliber of 58 mm of the skin of cellulose fibre to be produced was shaped into a tube and preimpregnated and coated on the inner and outer side with the dyed viscose through spinnerets of the type described in EP 0 267 489.

The tube was then coagulated in an acid spinning solution and regenerated to form cellulose hydrate gel and at the same time the leuco form of the indanthrene dyes contained therein was reconverted by oxidation with atmospheric oxygen into the original, insoluble dye, which adhered firmly to the cellulose material. The salts formed as a result of the precipitation and the acid taken up were then removed from the tube of regenerated cellulose in the subsequent wash. Prior to the final drying to a moisture content of about 7 wt.%, based on the total weight of the wrapping, the skin of cellulose fibre was passed through a conventional softening bath of glycerol.

Sausages of the salami type were produced using this skin of cellulose fibre. The sausage casing on the finally prepared sausage exhibited high transparency and an even dye.

Example 2

A skin of cellulose fibre having a nominal caliber of 50 mm was produced in a manner similar to that described in Example 1, but the sequence of the dye liquor preparation process was altered and the concentration of the dye and of the reducing agent was lowered.

In a sealable container of 30 litres in volume, 30.0 kg of a dye liquor was prepared in the following manner:-

3.105 kg	of a paste, consisting of	
3.0 kg	dye paste Red 179, C.I. 71130, pigment content approximately 22% (6300 Vocofil Rot, Ari Chemie) and	
0.105 kg	pigment paste black 7, C.I. 77266, pigment content approximately 35% (water-insoluble pigment)	

were weighed out. 10.455 kg of a 5 wt.% solution of carboxymethyl cellulose (CRT 30 GA, Wolff Walsrode) was added thereto.

A freshly prepared solution of 250 g sodium hydroxide solution (50 wt.%) and 300 g sodium dithionite in 2.5 kg water was added thereto, with gentle stirring with a paddle mixer, the rate of rotation being so adjusted that no air bubbles were introduced into the dye liquor.

After a residence time of 16 hours in the sealed container, this dye liquor was mixed continuously with viscose, in a ratio of 7.0 l dye liquor to 100 kg viscose. As described in Example 1, a tubular web of non-woven fabric was coated with the dyed viscose, regenerated and oxidised, then washed, passed through a softening bath and dried.

Sausages of the salami type were produced using this skin of cellulose fibre. The sausage casing on the finally prepared sausage exhibited high transparency and an even dye which, as regards the intensity, corresponded to the sausages in Example 1.

Example 3

A skin of cellulose fibre having a nominal caliber of 50 mm was produced in a manner similar to that described in the previous Examples, but here the dyeing was carried out using a brown vat dye. The sequence of the dye liquor preparation process and the concentration of the reducing agent corresponded to those in Example 2.

In a sealable container of 30 litres in volume, 30.0 kg of a dye liquor was prepared as follows:-

1.0 kg	of an indanthrene pigment paste of vat brown 57, having a pigment content of approximately 30% (223 Vocafil brown dye, Ari Chemie) was weighed out. To this was added
0.8364 kg	of a 5 wt.% solution of carboxymethyl cellulose (CRT 30 GA, Wolff Walsrode) and
12.196 kg	water. The incorporation of air was prevented by gentle stirring with a paddle mixer, during which a freshly prepared solution of
200 g	sodium hydroxide solution (50 wt.%) and
240 g	sodium dithionite
in 2.0 kg	water
	was added.

After a residence time of 16 hours in the sealed container, this dye liquor was mixed continuously with viscose, in a ratio of 6.2 l dye liquor to 100 kg viscose. A tubular web of non-woven fabric was coated with the dyed viscose, regenerated and oxidised, then washed, passed through a softening bath and dried, as described in Example 1. Sausages of the salami type were produced using this skin of cellulose fibre. The sausage casing on the finally prepared sausage exhibited high transparency of the even, reddish-brown dye.

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Claims

1. Process for producing dyed, tubular food wrappings from non-woven fabric coated with regenerated cellulose, characterised in that an alkaline dye liquor containing at least one dye which has been previously converted into an alkali-soluble form by chemical reduction and which can be converted into its insoluble form by oxidation is admixed to the viscose solution used for the production of the layer of regenerated cellulose, a tubular non-woven fabric is coated with the mixture of viscose solution and dye liquor, the viscose is coagulated and regenerated to form cellulose hydrate gel and the dye distributed in the viscose is reconverted into its insoluble form by oxidation.

2. Process according to claim 1, characterised in that dyes of the class of substances comprising the anthraquinone derivatives, preferably derivatives of anthrimidecarbazole, acylaminoanthraquinone, acridone, benzanthrone, violanthrone, isoviolanthrone, indanthrone, and derivatives of more highly condensed aromatic ring systems, preferably pyrenequinone, anthanthrone, flavanthrone, pyranthrone, perylenetetracarboxylic acid, naphthalene-tetracarboxylic acid as well as indigo derivatives and thioindigo derivatives are used.

3. Process according to claim 1 or 2, characterised in that for the chemical reduction of the coloured pigments, sodium dithionite or sodium sulfide is used in a quantity of 10 to 90 wt.%, preferably 20 to 80 wt.%, based on the pure, reducible dye in the dye liquor.

4. Process according to claims 1 to 3, characterised in that the addition of the dye liquor to the viscose is 2 to 26 litres, preferably 3 to 9 litres, per 100 kg viscose.

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5. Process according to claims 1 to 4, characterised in that the dye liquor contains in addition alkali- and acid-resistant non-reducible coloured pigments in a quantity of 3 to 12 wt.%, preferably 4 to 7 wt.%, based on the total quantity of dye and dye pigment.

6. Process according to claims 1 to 5, characterised in that the dye liquor contains cellulose ether, preferably carboxymethyl cellulose and/or methyl cellulose.

7. Tubular wrapping for food, in particular synthetic casing for sausages, made of non-woven fabric coated with regenerated cellulose, containing a transparent dye according to one of claims 1 to 6.

8. Tubular wrapping for food according to claim 7, characterised in that the outer or inner surface of the wrapping has a barrier layer which is impermeable to water vapour and oxygen.

9. Use of the tubular wrapping for food according to one of claims 7 or 8 as synthetic casings for raw sausage, sausages for boiling or sausages for cooking.

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Process for producing transparent dyed cellulose wrappings

Abstract

Process for producing dyed, tubular food wrappings from non-woven fabric coated with regenerated cellulose, characterised in that an alkaline dye liquor containing at least one dye which has been previously converted into an alkali-soluble form by chemical reduction and which can be converted into its insoluble form by oxidation is admixed to the viscose solution used for the production of the layer of regenerated cellulose, a tubular non-woven fabric is coated with the mixture of viscose solution and dye liquor, the viscose is coagulated and regenerated to form cellulose hydrate gel and the dye distributed in the viscose is reconverted into its insoluble form by oxidation; tubular food wrappings produced by this process and their use as synthetic casings for sausages.

COMBINED DECLARATION AND POWER OF ATTORNEY	ATTORNEY DOCKET NO

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name. I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought
on the invention entitled

METHOD FOR PRODUCING TRANSPARENT, COLOURED CELLULOSE SLEEVES

the specification of which is attached hereto,

or was filed on **April 16, 1999**

as a PCT Application Serial No. **PCT/EP99/02553**

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims.

I acknowledge the duty to disclose information which is material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s), the priority(ies) of which is/are to be claimed:

198 18 891.9	**Germany**	**April 28, 1998**
(Number)	(Country)	(Month/Day/Year Filed)

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose the material information as defined in Title 37, Code of Federal Regulations, §1.56 which occured between the filing date of the prior application and the national or PCT international filing date of this application:

(Application Serial No.)	(Filing Date)	(Status) (patented, pending, abandoned)
(Application Serial No.)	(Filing Date)	(Status) (patented, pending, abandoned)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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